(12) UK Patent Application (19) GB (11) 2 147 597 A

(43) Application published 15 May 1985

(21) Application No 8421431

(22) Date of filing 23 Aug 1984

(30) Priority data

(31) 8322643 8333462 (32) 23 Aug 1983 (33) GB 15 Dec 1983

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(51) INT CL4 C08L 61/28 C08K 3/34

(52) Domestic classification C3R 38C 38D2C 38D2D 38D3A C11 C12 C16 C17A C22 C23 C29 C5B1 C6X C8P L2B L2C1 L2CX L2X L4C L5C L6F V C3K 120 122 123 124 201 203 272 274 282 283 284 291 EF C3W 302 328 U1S 1362 3011 3015 C3K C3R

(56) Documents cited None

(58) Field of search C3R СЗК СЗМ

(54) Intumescent material

(57) An intumescent material comprises: one or more organic polyhydroxy compounds as carbonific; one or more organic polyamido compounds in free and jor combined form as spumific; an ammonium phosphate as activator for the carbonific; and an amino resin as binder, the material further comprising sodium silicate in an amount insufficient to prevent the material intumescing on being subjected to elevated temperature.

A method of manufacturing the intumescent material, comprises: preparing an aqueous paste or slurry comprising the ingredients of the material and allowing or causing the paste or slurry to harden.

SPECIFICATION

Intumesc nt mat rial

	and the state of t	5
5	The pr sent invention relates to intumescent mat rial. An intumescent material is a material which when heated or subjected to elevated tempera-	
		10
10		10
	. I I I I I I I I I I I I I I I I I I I	
	A second known type of intumescent material comprises components acting as soldering	
	in it is a labella de la compania d	
	The carbonific is a source of carbon char when the intumescent material is subjected to heat.	15
15	It is the principal constituent of the foam formed on intumescence. Mono-, di- and tri-	
	pentaerythritols, sugars and starch are examples of carbonifics. The spumific decomposes under fire conditions and releases gases which cause the carbon The spumific decomposes under fire conditions and releases gases which cause the carbon	
	The spumific decomposes under the contained and tributes grand guanidine, in free or char to expand. Polyamido compounds such as discyandiamide and guanidine, in free or	
	the different are exampled of OFGSDIC SDIFFIELD.	
20	The activator promotes the decomposition of the carbonitic. Antifolium phosphates dusit as	20
20		
	The binder may be an amine resin such as urga-termalgenyde resin of melanine-to-mar	
	mi i i i i i i i i i i i i i i i i i i	25
25	the amount of the process of the second of t	
	carbonific and spumific. Such a material needs to be formulated so as to exhibit an acidic reaction in water if it is to intumesce satisfactorily under fire conditions. This is because the	
	reaction in water it it is to intumesce satisfactoring direct in describing the decomposition of the ammonium phosphate has to yield phosphoric acid in order to promote the decomposition of the	
	,	
30	tables and motorials of the second type and comprising amino resin as binder are discussed	30
30	Bridge Barant and 1601131 Such Intilmescent Material Complises intigendictions	
	the decree sinked by a polyamido compound, the cross-linked resin acting as billed and	
	the relyamide compound acting as spumitic. The material also complises all alliforniting	
	the base of perioder and an organic polyhydroxy compound as carponing.	35
35	Intumescent materials, as described above, made from cross-linked amino resins, such as	-
	melamine-formaldehyde resin, may be regarded as heavily filled amino resins and hence are stronger and less friable than sodium-silicate based intumescent material.	
	re englications, intumescent materials based on amino resins, especially melanines	
	formaldehyde resignment he considered superior to intumescent materials comprising societies	
40	siliente in view of the shove-mentioned differences in their properties.	40
40	the same and the street costs with less than amino resilia autilia including	
	formaldehyde resin and hence intumescent materials based on sodium silicate can be manufac-	
	as the agest then intumescent materials based on amino resins.	
	The above-described intumescent materials comprising amino resins are generally manufac-	45
45	tured by making a thick aqueous slurry or paste of the required ingredients and allowing or	
	causing the slurry or paste to harden. In the case of intumescent materials as described in our aforementioned British patent, the	
	ability and a supply of paste is prepared from inter alla water-dispersible melamine-iornia-	
	dehyde resin, a polyamido compound such as dicyandiamide as cross-linking agent for the resin	
50	and an assemble and managementum phosphate.	50
	It would not be thought that a satisfactory incumescent material comprising components	
	period as earbonific, solumific and binder, ammonium phosphate acting as activator, and also	
	sodium silicate, could be manufactured at reduced cost (as compared with such a material	
	without sodium silicate) because the alkalinity of the sodium silicate would be expected to	55
55	prevent or hinder the yielding of phosphoric acid by the ammonium phosphate. Also in manufacturing the intumescent material, if sodium silicate were incorporated into the aqueous	
	slurry or paste, it would be expected to react with the resin, which gives an acidic reaction in	
	weeter, and provent the resin cross-linking properly.	
	We have now surprisingly discovered that a satisfactory and, in some respects superior,	
60	intumescent material may comprise a carbonific, an organic spumific, an amino resin, such as	60
	erosstinked melamine-formaldehyde resin, as binder, and sodium silicate.	
	Accordingly the present invention provides an intumescent material comprising; one or more	
	is polyhydromy compounds as carbonific; one or more organic polyamido compounds in	
	fre and/or combined form as spumific; an ammonium phosphate as activator for the carbonific;	65
65	and an amino resin as binder, the material further comprising sodium silicate in an amount	

insufficient to pr vent the material intumescing on being subjected to I vated t mp rature. It will be understood that the sodium silicat in a sinse constitutes a further spumific. The proportions of the organic polyhydroxy compound, the polyamido compound, the amm nium phosphate, the amino r sin, and any other ingredi nts (except sodium silicate) may 5 5 be varied as describ d in our British patent no. 1601131. The amino resin is preferably melamine-formaldehyde resin cross-linked by a polyamido compound, such as dicyandiamide or guanidine, which acts as spumific. The carbonific may conveniently be pentaerythritol. The activator is preferably monoammonium phosphate or may be ammonium polyphosphate. In order to provide satisfactory intumescence, the amount of sodium silicate preferably does 10 not exceed 12.7%, more preferably 11.5% by weight based on the total weight of the aforementioned carbonific, organic spumific (polyamido compound), activator and binder. Preferably the amount of sodium silicate does not exceed 32.6%, more preferably 29.6%, by weight based on the ammonium phosphate. 15 Preferably the amount of sodium silicate does not exceed 36%, more preferably 32.8%, by weight based on the amino resin excluding any cross-linkages thereof (e.g. cross-linking polyamido compound where the resin is melamine-formaldehye resin cross-linked by such polyamido compound). Preferably the amount of sodium silicate does not exceed 63.5%, more preferably 62.3%, by 20 20 weight based on the polyamido compound. Preferably the amount of sodium silicate does not exceed 171%, more preferably 155%, by weight based on the polyhydroxy compound. Preferably the amount of sodium silicate does not exceed 11%, more preferably 10%, by weight based on the ingredients of the material other than sodium silicate and any water 25 25 present. In order to provide a stiff "puff" as described below, the amount of sodium silicate preferably is at least 3.46%, more preferably 4.6%, by weight based on the total weight of the aforementioned carbonific, spumific (polyamido compound), activator and binder. Preferably the amount of sodium silicate is at least 8.9%, more preferably 11.9%, by weight 30 30 based on the ammonium phosphate. Preferably the amount of sodium silicate is at least 9.8%, more preferably 13.1% by weight based on on the amino resin excluding any cross-linkages thereof (e.g. cross-linking polyamido compound where the resin is melamine-formaldehyde resin cross-linked by such polyamido compound). Preferably the amount of sodium silicate is at least 18.7%, more preferably 24.9%, by weight based on the polyamido compound. Preferably the amount of sodium silicate is at least 46.7%, more preferably 62.3%, by weight based on the polyhydroxy compound. Preferably the amount of sodium silicate is at least 3%, more preferably 4%, by weight based 40 40 on the ingredients of the material other than sodium silicate and any water present. The intumescent material of the invention may further comprise polyvinyl acetate as additional binder to provide with the amino resin an interpenetrating polymer network system, thereby making the intumescent material less brittle. The intumescent material of the invention may further comprise a hardened setting agent such as hardened plaster of Paris. The intumescent material according to the invention has a number of advantages, apart from 45 its reduced cost, as compared with the second known type of intumescent material. On heating intumescent material according to the invention, intumescence occurs in two stages. A first stage of intumescence occurs at a relatively low temperature due to the presence of the sodium silicate and the second stage of intumescence occurs at a somewhat higher 50 50 temperature due to the ingredients of the second type of intumescent material. Moreover, the second stage intumescence tends to occur more slowly than if the sodium silicate were absent. Thus although the first stage intumescence may occur quickly, the second stage intumescence may be advantageously prolonged. Moreover the intumesced material or "puff" resulting from heating of intumescent material 55 according to the invention is stiff. In contrast the "puff" obtained from the conventional sodium 55 silicate-based intumescent material is hard and the "puff" obtained from intumescent material of the second type is soft and elastic. A stiff "puff" may be advantageous in certain situations for example in retarding warping of doors under fire conditions. Under such conditions a soft and elastic "puff" would deform easily 60 60 to accomodate the warping of the door and may slump to leave a gap between the intumesced material and the door whereas a hard "puff" would crack and crumble and may tend to force the door op n. However, a stiff "puff" has sufficient rigidity not to slump but do s not tend to crack and crumble r to force the door open. Mor over, when the intumescent material according to the invention intumesces through an 65 65 op ning in a hold r it tends to expand in a direction normal to the opening rather than

	spreading laterally from the opening and can thus seal larger gaps than the known intumescent						
	materials. The degree of "puff" or intumescence of the intumescent material may be up to 1/3 as large again as that of known intumescent materials of the second type and as described in our						
5	aforementioned British patent.	5					
	We have also found that the intumescent material according to the invention is not adversely affected by prolonged exposure to high concentrations of carbon dioxide or atmospheric						
	moisture. The material can withstand 100% relative humidity without its intumescent properties						
	being adversely affected.						
10		10					
	its intumescent potential well under accelerated ageing even for ageing equated with as much as ten years.						
	The intumescent material of the invention may be manufactured by preparing an aqueous						
	paste or slurry comprising: one or more organic polyhydroxy compounds as carbonific, one or						
15	more polyamido compounds as spumific, an ammonium phosphate as activator and an amino resin as binder, and sodium silicate, and allowing or causing the paste or slurry to harden, the	15					
	amount of sodium silicate being insufficient to prevent the material manufactured intumescing						
	on being subjected to elevated temperature.						
	Preferably the amino resin is water-dispersible melamine-formaldehyde resin, which becomes						
20	cross-linked by the polyamido compound. The polyamido compound may be dicyandiamide or guanidine, which can effect cross-linking of melamine-formaldehyde resin at normal ambient	20					
	temperature (15 to 25°C).						
	The amount of sodium silicate in relation to the other ingredients of the material is preferably						
	as specified above in relation to the intumescent material of the invention.	25					
25	The aqueous paste or slurry may further comprise water-dispersible polyvinyl acetate as binder (in addition to the amino resin) for the purpose referred to above and a setting agent such as	25					
	plaster of Paris.						
	Preferably the pH of the paste or slurry from which the intumescent material is prepared is not						
20	above 6.2, more preferably 6.1, otherwise the degree of intumescence of the material on	20					
30	heating may be inadequate. We have found that the paste or slurry runs very smoothly and is thus advantageous for	30					
	loading into holders in which it subsequently sets to form intumescent material. Also the paste						
	or slurry tends to be less adherent to the equipment used in its preparation than the aqueous						
25	mixture used in preparing the known intumescent materials and thus leaves the equipment	35					
33	cleaner. Preferably the pH of the paste or slurry is about 6.15 to provide an optimum combination of smoothness of the paste or slurry and adequate intumescence of the intumescent	33					
	material, although a lower pH would normally be chosen in order to provide greater intumes-						
	cence.						
40	The invention is illustrated by the following Examples: The sodium silicate used in the Examples was commercially available sodium silicate having	40					
40	an SiO : Na₂O molar ratio of 1:2. The plaster of Paris used in the Examples was of a kind which						
	exhibits high expansion during setting.						
	Example 1						
45	A powder mix of the following ingredients in the stated amounts was prepared:-	45					
	water-dispersible melamine formaldehyde resin powder 1900g						
	monoammonium phosphate 2100g						
50	dicyandiamide 1000g	50					
	pentaerythritol 400g						
	plaster of Paris 550g water-dispersible polyvinyl acetate powder 150g						
	water-dispersible polyvinyl acetate powder 150g wood flour (passing through 90 mesh) 125g						
55	Tool Annual Manager and Manager Annual Manager Annu	55					
	Total 6225g						

100 parts by weight of this powder mix were mixed with various amounts of sodium silicate powder, the resulting mixture worked up with water and the pH of the worked-up mixture 60 measured. The worked-up mixture was then allowed to set and dry. The set and dried mixture was then subject to elevated temperature to determine its degree of intumescence.

The results were as follows:

	•-										
	Amount of sodium										
	silicate (parts by										
5	weight)	pН	Intumescence	Remarks		5					
	4	< 6	good	stiff "puff" o	obtained						
10	6	< 6	very good	stiffer "	••	10					
10	8	∽ 6	11 17	stiffest "	••						
	10	6.1	good								
15	125	6.5	reduced			15					
	15	6.8	poor								
20	Example 2 The following composition was made as a dry powder										
25	water-dispersible melamine-formaldehyde resin monoammonium phosphate 2100g dicyandiamide 1000g pentaerythritol 400g plaster of Paris water-dispersible polyvinyl acetate powder 150g										
30	wood flour (passing through 96 mesh) 125g sodium silicate powder 498g										
35	Water was mixed with the above dry composition to form a thick aqueous slurry. The thick aqueous slurry was quickly poured into a tubular holder strip. The initial setting of the material started within 1–2 hours and chemical setting was well advanced after 2 days. The holder strip then slit along one face by a milling cutter and the holder with the hardened mixture was then kept at 40°C for 24 hours to mature the hardened mixture (i.e. to allow free water to evaporate and to complete the chemical setting). The holder containing the intumescent material is referred to herein as a first sealing device.										
40	A second sealing device was made in like manner to that described above but using no sodium silicate in preparing the aqueous slurry. It was found that the crack resistance of the intumescent material in the two holders was										
45	substantially identical. On heating the two sealing devices, the material therein intumesced, the material made using sodium silicate intumescing about ¹ / ₃ more than the other material. The intumesced mass provided by first sealing device (and comprising sodium silicate) was stiff and that provided by the second sealing device was soft and elastic.										
50	CLAIMS 1. An intumescent material comprising: one or more organic polyhydroxy compounds as carbonific; one or more organic polyamido compounds in free and/or combined form as spumific; an ammonium phosphate as activator for the carbonific; and an amino resin as binder, the material further comprising sodium silicate in an amount insufficient to prevent the material										
55	by a polyamido compound which acts as spumific. 4. A material according to any preceding claim, wherein the amount of sodium silicate is no										
60	more than 32.62% by weight based on the ammonium phosphate. 6. A material according to any preceding claim, wherein the amount of sodium silicate is no										
69	m re than 36% by weight based on the amino resinexcluding any cross-linkages there for an area of the second of th										

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8.	A materi	al according	to any	preceding	claim,	wher	in the	amount o	f sodium	silicate	does
not ex	ceed 119	6 by weight	based	n the tota	I weigh	nt of th	ne othe	r ingredie	ents of th	materi	al.

 A material according to any prec ding claim, wh rein the amount of s dium silicate is at least 3.46% by weight based on the total weight of the organic polyhydroxy compound, the polyamido compound, the ammonium phosphate and the amino resin.

10. A material according to any preceding claim, wherein the amount of sodium silicate is at least 8.9% by weight based on the ammonium phosphate.

11. A material according to any preceding claim, wherein the amount of sodium silicate is at least 9.8% by weight based on the amino resin excluding any cross-linkages thereof.

12. A material according to any preceding claim, wherein the amount of sodium silicate is at least 18.7% by weight based on the polyamino compound.

13. A material according to any preceding claim, wherein the amount of sodium silicate is at least 46.7% by weight based on the polyhydroxy compound.

14. A material according to any preceding claim, wherein the amount of sodium silicate is at 15 least 3% by weight based on the total weight of the other ingredients of the material.

15. A method of manufacturing an intumescent material, comprising: preparing an aqueous paste or slurry comprising one or more organic polyhydroxy compounds as carbonific, one or more polyamido compounds as spumific, an ammonium phosphate as activator and an amino resin as binder, and sodium silicate, and allowing or causing the paste or slurry to harden, the amount of sodium silicate being insufficient to prevent the material manufactured intumescing

20 amount of sodium silicate being insufficient to prevent the material manufactured intumescing on being subjected to elevated temperature.

16. A method according to claim 15, wherein the amino resin is water-dispersible melamine-formaldehyde resin and the aqueous paste or slurry comprises a polyamido compound as spumific and cross-linking agent for the resin.

17. A method according to claims 15 or 16, wherein the pH of the paste or slurry is not above 6.2.

Printed in the United Kingdom for Her Majesty's Stationery Office, Dd 8818935, 1985, 4235.
Published at The Patent Office, 25 Southempton Buildings, London, WC2A 1AY, from which copies may be obtained.